Climate Response and Radiative Forcing for Each Aerosol Species in CESM Prescribed from NCAR and Harvard Concentrations

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Model experiment setup

- CESM v1.0.3 with CAM4
- NCAR: use year 2000 aerosol emissions Harvard: use 2008 aerosol mass concentrations from GEOS-Chem
- Aerosol radiative forcing (ARF) is instantaneous radiative forcing estimated by calling the radiative transfer module twice.

Observation Comparisons - BC



Comparison with DC-8 ARCTAS observations



Aerosol Optical Depth





Dust: Optical Depth

NCAR tau







Black Carbon: Optical Depth





Dust: Radiative Forcing@ TOA



 0° 0° E 90° E 135° W 90° W 45° W -2° -4° -4° -6°

RF: radiative forcing

Black Carbon: Radiative Forcing@ TOA



RF: radiative forcing

90 Š

Radiative Forcing @ TOA in Arctic -- NCAR



Radiative Forcing @ TOA in Arctic -- Harvard



TREFHT: NCAR - Harvard



TREFHT: surface air temperature at reference height

TREFHT: NCAR - Harvard



TREFHT: surface air temperature at reference height

Summary

Harvard (or GEOS-Chem) concentrations for year 2008 has much higher aerosol concentrations than in NCAR concentrations for year 2000.

The RF for BC over the Sahara is in excess of 6 W/m², where NCAR is 1-2 W/m². In 2008 Harvard has >1 W/m² over the Arctic.

However, the global mean climate response is about the same for NCAR and Harvard data sets due to cancellation. But there are interesting hemispheric and regional differences.

The global mean climate differences are small in part because dust, sulfate, and OC effects oppose those from BC.

In the Arctic RF in the Harvard estimate is dominated by BC, but recall the Harvard estimate is for year 2008.

Spring BC Concentrations

Harvard data



Comparison with Arctic aerosol optical depth (AOD) observations

AOD across 8 Arctic AERONET Stations



OC is the main AOD component in summer due to large open fire sources

Absorption AOD and deposition of absorbing species (>65N)

Absorption = Mass cpt (g m⁻³) * Mass Absorption Efficiency (m² g⁻¹) [MAE BC=9.5m²g⁻¹, OC= $0.27m^2 g^{-1}$, Dust= $0.03m^2g^{-1}$]



Open Fires were the main source in spring and summer 2008 Significant contribution from non-BC aerosol in spring and summer to AAOD (27%) and Deposition (36%)

Decadal Trends Simulation 1980-2010

- MERRA meteorological fields at 4°x 5° horizontal resolution
- GFED3 open fire emissions for 1997 to 2010 [van der Werf et al., 2010]
- Anthropogenic BC and OC emission trends [Bond et al., 2007] with linear growth scaling factors for Russia and China for 2000-2008 (x2 in 2008)







TREFHT: NCAR - Harvard



RF: radiative forcing F_{TOA} : net flux at TOA TREFHT: surface air temperature at reference height